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10/014,519	12/14/2001	Betty Wu	19662-029001	3927
26161 75	10/16/2006		EXAMINER	
FISH & RICHARDSON PC			SINES, BRIAN J	
P.O. BOX 1022 MINNEAPOLIS, MN 55440-1022			ART UNIT PAPER NUM	
	•		1743	

DATE MAILED: 10/16/2006

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)				
Interview Summary	10/014,519	WU ET AL.				
merview dummary	Examiner	Art Unit				
	Brian J. Sines	1743				
All participants (applicant, applicant's representative, PTO	personnel):					
(1) Brian J. Sines.	(3)					
(2) <u>R. Bone</u> .	(4)					
Date of Interview: 11 October 2006.						
Type: a)⊠ Telephonic b)□ Video Conference c)□ Personal [copy given to: 1)□ applicant 2	2)  applicant's representative	e]				
Exhibit shown or demonstration conducted: d) Yes If Yes, brief description:	e)⊠ No.					
Claim(s) discussed: present claims.						
Identification of prior art discussed: <u>N/A</u> .						
Agreement with respect to the claims f)⊠ was reached. g	)□ was not reached. h)□ N	I/A.				
Substance of Interview including description of the general reached, or any other comments: <u>The non-final action, mailallowance</u> , mailed 10/2/2006. Typographical errors were concluded to include the second gas before final claim renumbering. Claim 46 was corrected to the second gas before final claim renumbering.	led 9/22/2006, is vacated due prected in the specification on actuator. Claim 21 was corre	to the subsequent notice of page 3 (see attached). ected to depend from claim 16				
(A fuller description, if necessary, and a copy of the amend allowable, if available, must be attached. Also, where no c allowable is available, a summary thereof must be attached	opy of the amendments that w					
THE FORMAL WRITTEN REPLY TO THE LAST OFFICE A INTERVIEW. (See MPEP Section 713.04). If a reply to the GIVEN A NON-EXTENDABLE PERIOD OF THE LONGER INTERVIEW DATE, OR THE MAILING DATE OF THIS INT FILE A STATEMENT OF THE SUBSTANCE OF THE INTE requirements on reverse side or on attached sheet.	last Office action has already OF ONE MONTH OR THIRTY ERVIEW SUMMARY FORM,	been filed, APPLICANT IS Y DAYS FROM THIS WHICHEVER IS LATER, TO				
Examiner Note: You must sign this form unless it is an	Bian	Jun				
Attachment to a signed Office action.	Examiner's sign	ature, if required				

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... IU !-... :... 200 material from the cells. These elements preferably operate differently from a valve, which would completely obstruct passage of material between upstream and downstream locations adjacent the valve. Rather, they typically provide resistance to fluid flow at a desired location (the lysing position) to thereby control fluid placement.

In one embodiment, the positioning element is disposed downstream of the lysing mechanism to position an upstream portion of a cell-containing sample (such as a microdroplet) in the lysing position. The positioning element preferably increases a surface tension of a downstream surface of the cell-containing sample to thereby inhibit downstream movement of the sample. For example, the positioning element may include an amount of reduced-wetting material, such as a hydrophobic material, disposed to contact a portion of the downstream surface of the cell-containing microdroplet.

In another embodiment, the positioning element is disposed upstream of the lysing zone to position a downstream portion of the cell-containing microdroplet in the lysing position. The positioning element includes a vent, which substantially equalizes a gas pressure upstream of the cell-containing microdroplet with a gas pressure downstream of the cell-containing microdroplet to thereby stop downstream movement of the cell-containing microdroplet. When the microdroplet is in the lysing position. A valve is preferably disposed to subsequent obstruct passage of gas between the lysing zone and the vent to allow an upstream gas pressure to once again move the droplet further downstream for additional processing. For example, the microfluidic system may include a mixing zone downstream of the enrichment zone and/ or lysing zone, to mix the microdroplet which emerges from these zones with a predetermined amount of reagent material.

In another aspect, the invention relates to a microfluidic substrate for processing the intracellular contents of cells suspended in fluids. The substrate includes a lysing module, a microdroplet formation module, mixing module and an amplification module. The lysing module releases intracellular material from cells within the sample to thereby form a lyced sample. The microdroplet formation module then forms a first microdroplet of fluid from the lysed sample and forwards it to a mixing module for mixing with a microdroplet of reagent. The amplification module amplifies intercellular material within the microdroplet formed from the mixture.

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